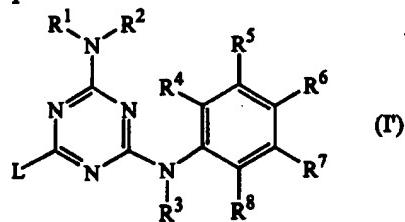


In the Claims:

Add new independent claim 19, as in the following clean list of claims:

1. (Five times amended) A compound of formula



a pharmaceutically acceptable acid addition salt or a stereochemically isomeric form thereof, wherein

R¹ and R² are each independently selected from hydrogen; hydroxy; amino; C₁-alkyl; C₁-alkyloxy; C₁-alkylcarbonyl; C₁-alkyloxycarbonyl; Ar¹; mono- or di(C₁-alkyl)amino; mono- or di(C₁-alkyl)aminocarbonyl; dihydro-2(3H)-furanone; C₁-alkyl substituted with one or two substituents each independently selected from amino, imino, aminocarbonyl, aminocarbonylamino, hydroxy, hydroxyC₁-alkyloxy, carboxyl, mono- or di(C₁-alkyl)amino, C₁-alkyloxycarbonyl and thienyl; or

R¹ and R² taken together may form pyrrolidinyl, piperidinyl, morpholinyl, azido or mono- or di(C₁-alkyl)aminoC₁-alkylidene;

R³ is hydrogen, Ar¹, C₁-alkylcarbonyl, C₁-alkyl, C₁-alkyloxycarbonyl, C₁-alkyl substituted with C₁-alkyloxycarbonyl; and

R⁴, R⁵, R⁷ and R⁸ are each independently selected from hydrogen, hydroxy, halo, C₁-alkyl, C₁-alkyloxy, cyano, aminocarbonyl, nitro, amino, trihalomethyl or trihalomethoxy;

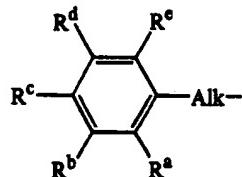
R⁶ is aminocarbonyl;

L is C₁-₁₀alkyl; C₃-₁₀alkenyl; C₃-₁₀alkynyl; C₃-₇cycloalkyl; or

L is C₁-₁₀alkyl substituted with one or two substituents independently selected from the group consisting of C₃-₇cycloalkyl; indolyl or indolyl substituted with one, two, three or four substituents each independently selected from halo, C₁-alkyl, C₁-alkyloxy, cyano, aminocarbonyl, nitro, amino, trihalomethyl, trihalomethoxy, or C₁-alkylcarbonyl; and phenyl or phenyl substituted with one, two, three, four or five substituents each independently selected from halo, hydroxy, C₁-alkyl, C₁-alkyloxy, cyano, aminocarbonyl, nitro, amino, trihalomethyl, trihalomethoxy, or C₁-alkylcarbonyl; and,

Ar^1 is phenyl, or phenyl substituted with one, two or three substituents each independently selected from halo, C_{1-6} alkyl, C_{1-6} alkyloxy, cyano, nitro or trifluoromethyl.

2. (Once Amended) A compound according to claim 1 wherein R^1 and R^2 are each independently selected from hydrogen, C_{1-6} alkyl, Ar^1 or mono- or di(C_{1-6} alkyl)aminocarbonyl; or R^1 and R^2 taken together may form pyrrolidinyl, piperidinyl or morpholinyl; R^3 is hydrogen, C_{1-6} alkyl or Ar^1 ; and Ar^1 is phenyl, or phenyl substituted with one, two or three substituents each independently selected from halo, C_{1-6} alkyl, C_{1-6} alkyloxy, cyano, nitro or trifluoromethyl; and L is a radical of formula



wherein Alk is C_{1-6} alkanediyl;

R^a , R^b , R^c , R^d , R^e , R^4 , R^5 , R^7 and R^8 are each independently selected from hydrogen, halo, C_{1-6} alkyl, C_{1-6} alkyloxy, cyano, aminocarbonyl, nitro, amino, trihalomethyl or trihalomethoxy; or

R^a and R^b taken together may form a bivalent radical of formula

-CH=CH-NR⁹. (a-1),

-NR⁹-CH=CH-. (a-2),

wherein R^9 is hydrogen or C_{1-4} alkyl.

3. (Twice amended) A compound according to claim 1 wherein L is C_{3-10} alkenyl or C_{1-2} alkyl substituted with one or two substituents independently selected from C_{3-7} cycloalkyl; indolyl or indolyl substituted with one, two, three or four substituents each independently selected from halo, C_{1-6} alkyl, C_{1-6} alkyloxy, cyano, aminocarbonyl, nitro, amino, trihalomethyl, trihalomethoxy, C_{1-6} alkylcarbonyl; phenyl or phenyl substituted with one, two, three, four or five substituents each independently selected from halo, hydroxy, C_{1-6} alkyl, C_{1-6} alkyloxy, cyano, aminocarbonyl, nitro, amino, trihalomethyl, trihalomethoxy, C_{1-6} alkylcarbonyl.

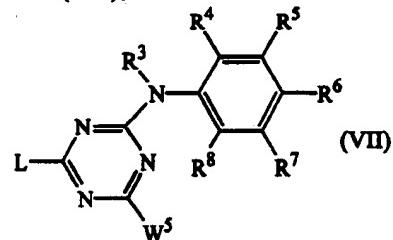
4. (Once Amended) A compound according to claim 3 wherein L is 2,6-dichlorophenylmethyl.

6. (Once Amended) A compound according to claim 4 wherein NR¹R² is other than amino.

11. (Once Amended) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a therapeutically active amount of a compound as claimed claim 1.

18 (Unchanged) A method of treating a subject suffering from HIV (Human Immunodeficiency Virus) infection comprising administering to the subject a therapeutically effective amount of the compound of claim 1.

19. (New) A compound of formula (VII),



wherein

R¹ and R² are each independently selected from hydrogen; hydroxy; amino; C₁₋₆alkyl; C₁₋₆alkyloxy; C₁₋₆alkylcarbonyl; C₁₋₆alkyloxycarbonyl; Ar¹; mono- or di(C₁₋₆alkyl)amino; mono- or di(C₁₋₆alkyl)aminocarbonyl; dihydro-2(3H)-furanone; C₁₋₆alkyl substituted with one or two substituents each independently selected from amino, imino, aminocarbonyl, aminocarbonylamino, hydroxy, hydroxyC₁₋₆alkyloxy, carboxyl, mono- or di(C₁₋₆alkyl)amino, C₁₋₆alkyloxycarbonyl and thienyl; or

R¹ and R² taken together may form pyrrolidinyl, piperidinyl, morpholinyl, azido or mono- or di(C₁₋₆alkyl)aminoC₁₋₄alkylidene;

R³ is hydrogen, Ar¹, C₁₋₆alkylcarbonyl, C₁₋₆alkyl, C₁₋₆alkyloxycarbonyl, C₁₋₆alkyl substituted with C₁₋₆alkyloxycarbonyl; and

R⁴, R⁵, R⁷ and R⁸ are each independently selected from hydrogen, hydroxy, halo, C₁₋₆alkyl, C₁₋₆alkyloxy, cyano, aminocarbonyl, nitro, amino, trihalomethyl or trihalomethoxy;

R⁶ is aminocarbonyl;

W⁵ is halo;

L is C₁₋₁₀alkyl; C₃₋₁₀alkenyl; C₃₋₁₀alkynyl; C₃₋₇cycloalkyl; or

L is C₁₋₁₀alkyl substituted with one or two substituents independently selected from the group consisting of C₃₋₇cycloalkyl; indolyl or indolyl substituted with one, two, three or four substituents each independently selected from halo, C₁₋₆alkyl, C₁₋₆alkyloxy, cyano, aminocarbonyl, nitro, amino, trihalomethyl, trihalomethoxy, or C₁₋₆alkylcarbonyl; and phenyl or phenyl substituted with one, two, three, four or five substituents each independently selected from halo, hydroxy, C₁₋₆alkyl, C₁₋₆alkyloxy, cyano, aminocarbonyl, nitro, amino, trihalomethyl, trihalomethoxy, or C₁₋₆alkylcarbonyl; and, Ar¹ is phenyl, or phenyl substituted with one, two or three substituents each independently selected from halo, C₁₋₆alkyl, C₁₋₆alkyloxy, cyano, nitro or trifluoromethyl.